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(54) Title: BISPECIFIC ANTISENSE OLIGNUCLEOTIDES THAT INHIBIT IGFBP-2 AND IGFBP-5 AND METHODS OF USING SAME

(57) Abstract: Bispecific antisense oligonucleotides which consist essentially of a sequence of bases that is complementary to portions of both the gene encoding human IGFBP-2 and the gene encoding human IGFBP-5 are useful in as antisense therapeutics in the treatment of endocrine-regulated cancers.

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BISPECIFIC ANTISENSE OLIGNUCLEOTIDES THAT
INHIBIT IGFBP-2 AND IGFBP-5 AND METHODS OF USING SAME

Background of the Invention

This present application relates to bispecific antisense oligonucleotides that inhibit IGFBP-2 and IGFBP-5 and methods of using same in the treatment of endocrine-regulated tumors (for example, breast, prostate, ovarian and colon cancers).

5 Prostate cancer is the most common cancer that affects men, and the second leading cause of cancer deaths in men in the Western world. Because prostate cancer is an androgen-sensitive tumor, androgen withdrawal, for example via castration, is utilized in some therapeutic regimens for patients with advanced prostate cancer. Androgen withdrawal leads to extensive apoptosis in the prostate tumor, and hence to a regression of the disease. However, castration-induced apoptosis is not complete, and a progression of surviving tumor cells to androgen-independence ultimately occurs. This progression is the main obstacle to improving survival and quality of life, and efforts have therefore been made to target androgen-independent cells. These efforts have focused on non-hormonal therapies targeted against androgen-independent tumor cells, however as of a 1998 report, no non-hormonal agent had improved survival. Oh et al., *J. Urol* 160: 1220-1229 (1998) Alternative approaches are therefore indicated.

15 Insulin-like growth factor (IGF)-I and IGF-II are potent mitogens for many normal and malignant cells. Accumulating evidence suggests that IGFs play an important role in the pathophysiology of prostatic disease and breast cancer. Boudon et al., *J. Clin. Endocrin. Metab.* 81: 612-617 (1996); Angelloz-Nicoud et al., *Endocrinology* 136: 5485-5492 (1995); Nickerson et al., *Endocrinology* 139: 807-810 (1998); Figueroa et al., *J. Urol.* 159: 1379-1383 (1998).

20 The biological response to IGF's is regulated by various factors, including IGFBPs. To date, six IGFBPs have been identified whose function is believed to involve modulation of the biological actions of the IGFs through high affinity interactions. Rajaram et al., *Endocrin. Rev.* 18: 801-813 (1997). However, some evidence suggests biological activity for IGFBPs that are independent of IGFs, *Id.*, Andress et al., *J. Biol. Chem.* 267: 22467-22472 (1992); Oh et al., *J. Biol. Chem.*

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268: 14964-14971 (1993), and both stimulatory and inhibitory effects of IGFBPs on cell proliferation have been reported under various experimental conditions. Andress et al., *supra*; Elgin et al., *Proc. Nat'l. Acad. Sci. (USA)*, 84, 3254-3258 (1987); Huynh et al., *J. Biol. Chem.* 271: 1016-1021 (1996); Damon et al., *Endocrinology* 139: 3456-3464 (1998). Thus, the precise function role of IGFBPs remains controversial. Because of this, while the reported results implicate IGF in prostate and breast cancer, they do not clearly suggest a therapeutic approach based upon this involvement.

PCT Publication WO 01/05435, which is incorporated herein by reference, describes a method for treating hormone-regulated tumors (for example, breast and prostatic tumors) in mammals, including humans, by administration of an antisense oligodeoxynucleotide which is complementary to a portion of the gene encoding IGFBP-5. PCT Publication No. WO 02/22642, which is incorporated herein by reference, describes a method are provided for the treatment of prostate and other endocrine tumors in mammals, including humans, by administration of an antisense oligodeoxynucleotide which is complementary to a portion of the gene encoding IGFBP-2.

The present invention utilizes bispecific antisense oligodeoxynucleotides targeted to both IBFBP-2 and IGFBP-5 as a treatment for endocrine-regulated cancers. Antisense oligodeoxynucleotides are chemically modified stretches of single-stranded DNA that are complementary to mRNA regions of a target gene, and thereby effectively inhibit gene expression by forming RNA/DNA duplexes. Figueroa, et al., *J. Urol.*, 159: 1379-1383 (1998). Phosphorothioate oligodeoxynucleotides are stabilized to resist nuclease digestion by substituting one of the nonbridging phosphoryl oxygen of DNA with a sulfur. Recently, several antisense oligodeoxynucleotides specifically targeted against genes involved in neoplastic progression have been evaluated both *in vitro* and *in vivo*, and demonstrated the efficacy of antisense strategy as potential therapeutic agents. Monia, et al. *Nature Med.* 2: 668-675 (1996.); Cucco, et al., *Cancer Res.* 56: 4332-4337 (1996); Ziegler, et al., *J. Natl. Cancer Inst.* 89: 1027-1036 (1997); Jansen, et al., *Nature Med.* 4: 232-234 (1998).

Summary of the Invention

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The present invention provides a method for treating endocrine-regulated tumors (for example, breast, prostate, ovarian and colon cancers) in mammals, including humans, by administration of a bispecific antisense oligodeoxynucleotide which is complementary to portions of both the gene encoding IGFBP-2 and the gene encoding IGFBP-5. The administration of such an oligodeoxynucleotide was shown to reduce levels of both IGFBP-2 and IGFBP-5 in PC3 and LNCaP cells, and the reduction of expression of the proteins has been previously shown to reduce proliferation of tumor cells, and also to delay the progression to androgen independence. Thus, in accordance with the invention we provide methods for treatment of prostate cancer in mammals, including humans, and for delaying the progression of prostate tumors to androgen independence comprising the step of administering to the mammal a therapeutically effective amount of a bispecific antisense oligodeoxynucleotide which is complementary to portions of both the nucleic acid sequence encoding IGFBP-2 and the nucleic acid sequence encoding IGFBP-5, and which hybridizes with such sequences to inhibit or otherwise reduce the expression of IGFBP-2 and IGFBP-5.

The application further provides bi-specific antisense sequences that can be used in the method of the invention.

Brief Description of the Figures

Fig. 1 the amount of IGFBP-2 and IGFBP-5 expression observed in LNCaP and PC3 cells, respectively, upon administration of the three bispecific antisense oligonucleotide, a mismatch control (MM), or no oligonucleotide (control).

Fig. 2 shows the increase in IGFBP-2 expression in prostate cancer cells during progression to androgen independence.

Figs. 3A-E shows inhibition of IGFBP-2 and 5 in prostate cancer and bone cells using various antisense oligonucleotides.

Figs. 4 and 5 show real time PCR results for levels of IGFBP-2 and IGFBP-5 in RT4 bladder cancer cells after treatment with antisense oligonucleotides of the invention.

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Fig. 6 shows results of real time PCR measurement of IGFBP-5 in MSF human fetal fibroblast cells after antisense treatment.

Fig. 7 shows the results of real time PCR measurements of IGFBP-2 and IGFBP-5 LNCaP/msf in mixed tumors and in intact fetal bone fragments

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Detailed Description of the Invention

The present invention provides bispecific antisense oligonucleotides which consist essentially of a sequence of bases that is complementary to portions of both the gene encoding IGFBP-2 and the gene encoding IGFBP-5, and that is sufficient length to act as an inhibitor of the effective amount of IGFBP-2 and IGFBP-5 (in general at least 15 bases). As used in the specification and claims of this application, this language means that substantially all of the antisense oligonucleotide is complementary to a portion of each gene sequence. The invention does not, however, exclude minor modifications in sequence, such as the addition of one or two terminal bases, or single base substitutions which might depart from perfect complementarity but which still function as an inhibitor of the effective amount of IGFBP-2 and IGFBP-5.

The phrase "inhibitor of the effective amount" takes into account the fact that antisense oligonucleotides may function by different mechanisms. The effective amount of IGFBP-2 or IGFBP-5 is the amount that is present in a functional state in the cell. Reduction of this amount by administration of antisense oligonucleotides may occur through restricting production of the IGFBP (at the transcription or translation level) or by degrading the IGFBP at a rate faster than it is being produced. Further, it will be appreciated that the inhibition which is referred to is one which occurs when the IGFBP would otherwise be present if the antisense oligonucleotide had not been administered. As discussed below, not all cells that are potential therapeutic targets express both IGFBP-2 and IGFBP-5.

Specific antisense oligonucleotides according to the invention consist essentially of a series of bases as set forth in Seq. ID. No. 1 through 7. These sequences are set forth in Table 1.

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Table 1	
Seq. ID NO.	Sequence
1	ggtgtagacgccgcacg
2	gcagcgagcccctgg
3	gcagcagccgcagcccggctcc
4	agccgcagcccggctcct
5	cagcagccgcagcccggctc
6	gcagcagccgcagcccggct
7	agcagccgcagcccggctcc

The antisense oligonucleotides employed may be modified to increase the stability of the antisense oligonucleotide *in vivo*. For example, the antisense oligonucleotides may be employed as phosphorothioate derivatives (replacement of a non-bridging phosphoryl oxygen atom with a sulfur atom) which have increased resistance to nuclease digestion. Increased antisense oligonucleotide stability can also be achieved using molecules with 2-methoxyethyl (MOE) substituted backbones as described generally in US Patent No. 6,451,991 and US Patent Application No. 10/080,794 which are incorporated herein by reference.

The compositions of the present invention can be used for treatment of endocrine-regulated tumors (for example, breast, prostate, ovarian and colon cancers) in mammals, including humans, by administration of a bispecific antisense oligonucleotide in accordance with the invention. Administration of antisense oligonucleotides can be carried out using the various mechanisms known in the art, including naked administration and administration in pharmaceutically acceptable carriers. For example, lipid carriers for antisense delivery are described in US Patents Nos. 5,855,911 and 5,417,978 which are incorporated herein by reference. In general, the antisense is administered by intravenous, intraperitoneal, subcutaneous or oral routes. Where the oligonucleotides are administered in a pharmaceutically acceptable carrier, the carrier is generally free from substances which produce toxic or other harmful reactions when administered to humans.

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The amount of antisense oligonucleotide administered is one effective to reduce the effective amount of levels of IGFBP-2 and/or IGFBP-5 in the endocrine-regulated tumor cell of concern. As noted above, in the context of the present invention, applicants do not intend to be bound by any specific mechanism by which this reduction may occur, although it is noted that the reduction may occur as a result of reduced expression of IGFBP-2 and -5 if the antisense molecule interferes with translation of the mRNA, or via an RNase mediated mechanism. Furthermore, it will be appreciated that the appropriate therapeutic amount will vary both with the effectiveness of the specific antisense oligonucleotide employed, and with the nature of any carrier used. The determination of appropriate amounts for any given composition is within the skill in the art, through standard series of tests designed to assess appropriate therapeutic levels.

The method for treating endocrine-regulated cancer in accordance with the invention may further include administration of chemotherapy agents and/or additional antisense oligonucleotides directed at different targets. For example, conventional chemotherapy agents such as taxol (paclitaxel or docetaxel) and mitoxanthrone may be used. Similarly, combinations of the bispecific antisense oligonucleotide of the invention with other antisense sequences such as antisense Bcl-2 oligonucleotide, TRPM-2 (clusterin) oligonucleotide, IGFBP-2 or IGFBP-5 oligonucleotide may be used.

The effectiveness of the present invention is founded on solid theoretical and experimental bases. Activation of alternative growth factor pathways following androgen withdrawal is one mechanism mediating androgen independent (AI) progression in advanced prostate cancer. Insulin-like growth factor-I (IGF-I) activation is modulated by a family of insulin-like growth factor binding proteins (IGFBPs). Although IGFBP-2 and IGFBP-5 are among the most commonly over-expressed genes in hormone refractory prostate cancer, the functional significance of changes in IGF-I signaling during AI progression remain poorly defined. AI progression of prostate cancer leads to treatment resistance and ultimately culminates in the development of bone metastases, the most ominous sign of clinical progression in prostate cancer patients with advanced disease. This development usually precedes death by 12 - 18

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months. The bone environment, particularly rich in IGFBP-5, has long been identified as an attractive "soil" for supporting prostate cancer metastasis. The crucial IGF-1 signaling components IGFBP-2 and IGFBP-5 may not only be key to AI progression but also to the site-specific metastasis of prostate cancer. In an attempt to address these issues, we characterized changes in IGFBP-2 and IGFBP-5 in prostate cancer models after androgen withdrawal and evaluated their functional significance in AI progression using gain-of-function and loss-of-function analyses.

IGFBP-2 mRNA and protein levels increase 2-3-fold after androgen withdrawal in LNCaP cells in vitro in LNCaP tumors during AI progression *in vivo*. Increased IGFBP-2 levels after castration were also identified using a human prostate tissue microarray of untreated and post-hormone therapy-treated prostatectomy specimen. LNCaP cell transfectants (LNBP-2) that stably over-expressed IGFBP-2 had a shorter cell doubling-time and a lower rate of apoptosis in the absence of androgens. Reporter assays demonstrated that this increased growth potential did not result from IGFBP-2-mediated transcription of androgen-response elements. *In vivo*, the LNBP-2 cell line formed significantly more tumors in castrate mice and progressed to androgen independence more rapidly compared with a control cell line. Antisense oligonucleotides targeting IGFBP-2 reduced IGFBP-2 mRNA and protein expression by >70% in a dose-dependent and sequence-specific manner. Antisense oligonucleotide-induced decreases in IGFBP-2 reduced LNCaP cell growth rates and increased apoptosis 3-fold. LNCaP tumor growth and serum PSA levels in mice treated with castration plus adjuvant IGFBP-2 antisense oligonucleotides were significantly reduced compared to mismatch control oligonucleotides.

Similar findings were obtained with gain-of-function and loss-of-function analyses of IGFBP-5 in IGFBP-5 expressing prostate cancer models. The growth rates of IGFBP-5 transfected LNCaP cells were significantly faster compared to either the parental or vector-only transfected LNCaP cells in both the presence and absence of dihydrotestosterone. IGFBP-5-induced increases in LNCaP cell proliferation occurs through both IGF-I-dependent and -independent pathways, with corresponding increases in the cyclin D1 mRNA expression and the fraction of cells in S + G2/M phases of the cell cycle. Changes in Akt/protein kinase B (PKB), a

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downstream component of phosphatidylinositol 3'-kinase (PI3K) pathway, in the LNCaP sublines also paralleled changes in their growth rates. Although treatment with a PI3K inhibitor induced apoptosis in both control and IGFBP-5-overexpressing LNCaP cells, this PI3K inhibitor-induced apoptosis was prevented by exogenous IGF-I treatment only in IGFBP-5 transfectants; suggesting that IGFBP-5 overexpression can potentiate the antiapoptotic effects of IGF-I. Furthermore, tumor growth and serum PSA levels increased several fold faster in mice bearing IGFBP-5-transfected LNCaP tumors after castration despite having similar tumor incidence and tumor growth rates with controls when grown in intact mice before castration. These data suggest that IGFBP-5 overexpression in prostate cancer cells after castration is an adaptive cell survival mechanism that helps potentiate the antiapoptotic and mitogenic effects of IGF-I, thereby accelerating progression to androgen-independence through activation of the PI3K-Akt/PKB signaling pathway. Systemic administration of IGFBP-5 antisense oligonucleotides significantly delayed time to progression to androgen independence and inhibited growth of AI recurrent tumors. IGFBP-5 expression in human bone was confirmed.

Inhibiting IGFBP-2 and IGFBP-5 expression using targeted antisense technologies provides a treatment platform to delay AI progression and bone metastasis in prostate cancer patients and patients with other malignancies expressing these key regulators of IGF-1 signaling. By targeting both IGFBP-2 and IGFBP-5 simultaneously with a single antisense oligonucleotide the complications associated with use of a cocktail of antisense oligonucleotides targeting either of these compounds separately are avoided.

The application is further described in the following non-limiting examples.

EXAMPLE 1

LNCaP cells were treated with treated with 500 nM concentrations of antisense oligonucleotides of Seq. ID Nos. 1, 2 or 3 or a mismatch control. Levels of IGFBP-2 were measured. The result are summarized in Fig. 1A.

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PC3 cells were treated with treated with 500 nM concentrations of antisense oligonucleotides of Seq. ID Nos. 1, 2 or 3 or a mismatch control. Levels of IGFBP-5 were measured. The result are summarized in Fig. 1B.

5

EXAMPLE 2

Samples of human prostate cancer cells were sectioned and immunostained to detect IGFBP-2. Tissue microarray scoring showed expression of IGFBP-2 was essentially absent in benign, non-cancerous tissues but increased with time to highest levels after androgen independent was achieved. (Fig. 2)

10

EXAMPLE 3

Osseous metastases of human prostate cancer was evaluated in 15 specimens. Human prostate cancer expressed IGFBP-2 in very high levels, but also expressed IGFBP-5. Human bone osteocytes expressed high levels of IGFBP-5 and lower levels of IGFBP-2.

15

EXAMPLE 4

Bispecific antisense oligonucleotides were used to treat PC3, LNCaP and bone cells at concentration of 500 nM, and the amount of inhibition of IGFBP-2 or IGFBP-5 was measured using real time PCR. Oligonucleotides of Seq. ID Nos. 2, 4, 5 and 7 were tested, and all were effective to reduce the detected amount of the IGFBP measured. (Fig. 3A) Reductions of up to 70% in IGFBP-2 levels were also observed in A549 lung cells using 500 nM of Seq. ID Nos. 4 and 5. (Fig. 3B) Seq. ID No. 4 (500 nM) was also shown to be effective to inhibit cell growth of LNCaP cells and reduce the cell number by more than 90%. (Fig. 3C) Figs. 3D and E respectively show results for inhibition of IGFBP-5 levels in PC3 cells with 500nM of Seq ID Nos. 1, 2, 4 5, 6 and 7; and inhibition of IGBFP-5 levels in human fetal bone fobroblast cells with 500 nM of Seq. ID Nos. 1, 2, 4, 5, 6 and 7.

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EXAMPLE 5

Real time PCR was used to measure the amount IGFBP-5 in MSF human fetal fibroblast cells after treatment with antisense oligonucleotide and LIPOFECTIN (4 µg/ml). Cells were plated *in vitro* and treated with a four-hour pulse of 500 nM oligonucleotide followed by a 20 hour period in normal medium plus 5% serum. A second four-hour pulse was repeated on day 2 and the cells were counted on day 3. The most active antisense oligonucleotides were Seq. ID Nos. 4 and 5. (Fig. 6)

EXAMPLE 6

Real time PCR was used to evaluate the amounts of IGFBP-2 and IGFBP-5 in human bladder cancer (RT4) following treatment with varying amounts of antisense oligonucleotides (Seq. ID Nos. 4 and 5) and 4 µg/ml LIPOFECTIN. As shown in Figs. 4 and 5, a dose dependent response was observed to both antisense oligonucleotides at concentrations ranging from 50 to 500 nM.

EXAMPLE 7

Real time PCR was used to measure IGFBP-2 and IGFBP-5 in LNCaP/msf mixed tumors and in intact fetal bone fragments. The results are shown in Fig. 7.

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CLAIMS

1. A bispecific antisense oligonucleotide which consist essentially of a sequence of bases that is complementary to portions of both the gene encoding IGFBP-2 and the gene encoding IGFBP-5, and which is sufficient length to act as an inhibitor of the effective amount of IGFBP-2 and IGFBP-5.
2. The antisense oligonucleotide according to claim 1, wherein the oligonucleotide consists essentially of a series of bases as set forth in Seq. ID. No. 1.
3. The antisense oligonucleotide according to claim 1, wherein the oligonucleotide consists essentially of a series of bases as set forth in Seq. ID. No. 2.
4. The antisense oligonucleotide according to claim 1, wherein the oligonucleotide consists essentially of a series of bases as set forth in any of Seq. ID. Nos. 3 through 7.
5. A method for making a pharmaceutical composition for treatment of endocrine-regulated cancers, comprising the steps of combining a bispecific antisense oligonucleotide in accordance with any of claims 1-4 with a pharmaceutically acceptable carrier for intravenous, intraperitoneal, subcutaneous or oral administration.
6. A method for treating an endocrine-regulated cancer in a subject suffering from the endocrine-regulated cancer, comprising the step of administering to the subject a bispecific antisense oligonucleotide in accordance with any of claims 1-4 in an amount effective to reduce the effective levels of IGFBP-2 and/or IGFBP-5 in cells of the endocrine-regulated cancer.
7. The method of claim 6, wherein the cancer is prostate cancer.
8. The method of claim 6, wherein the cancer is breast cancer.

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9. A pharmaceutical composition comprising an antisense oligonucleotide in accordance with any of claims 1-4 and a pharmaceutically acceptable carrier.

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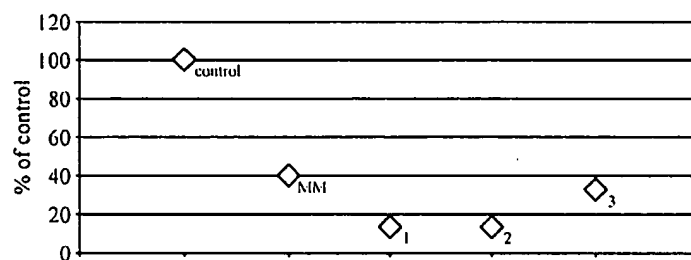


FIG. 1A

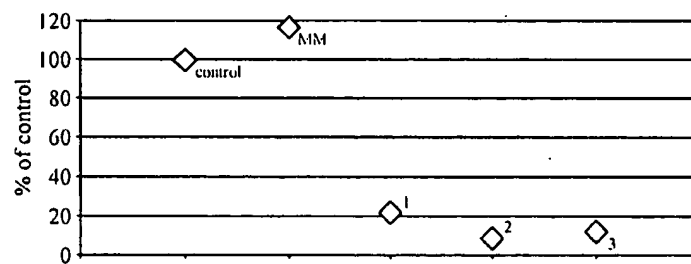


FIG. 1B

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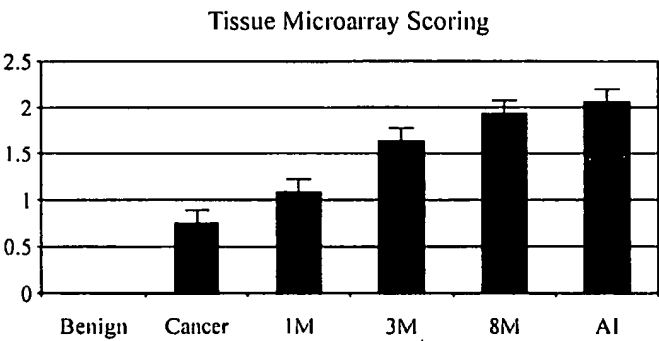


FIG. 2

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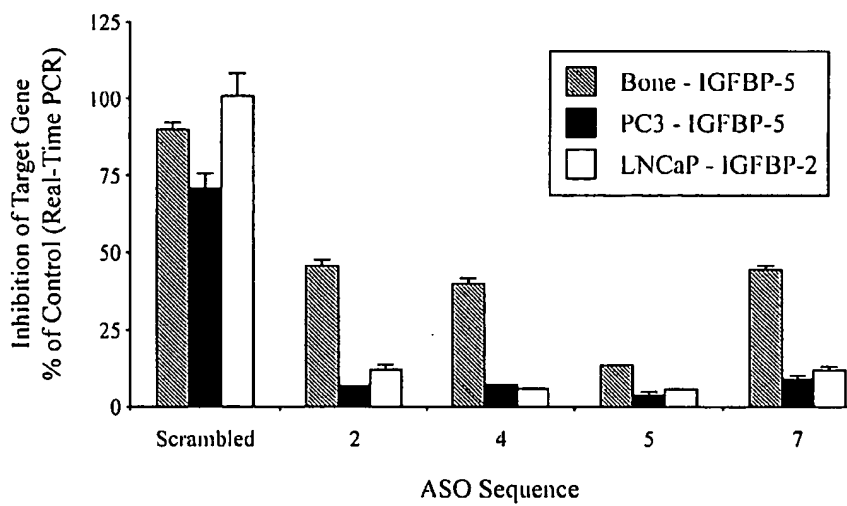


FIG. 3A

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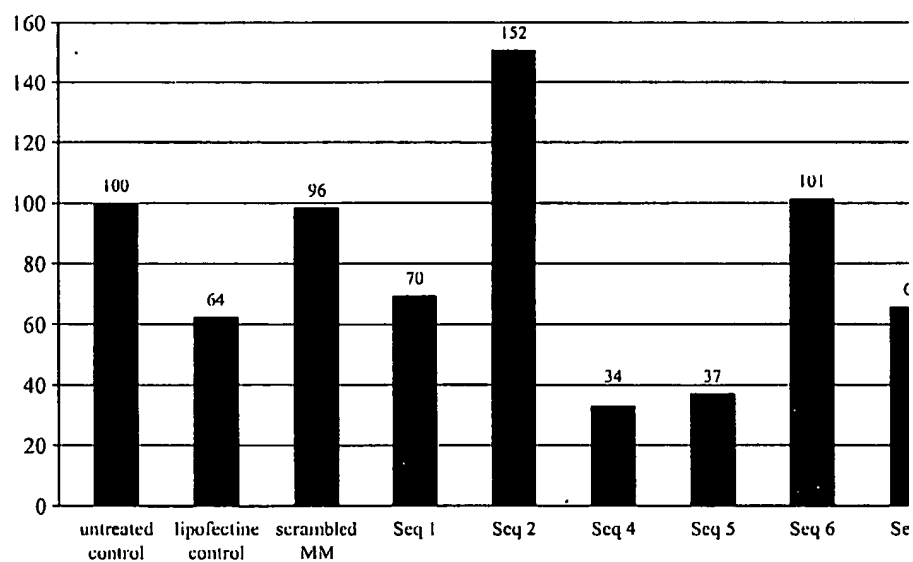


FIG. 3B

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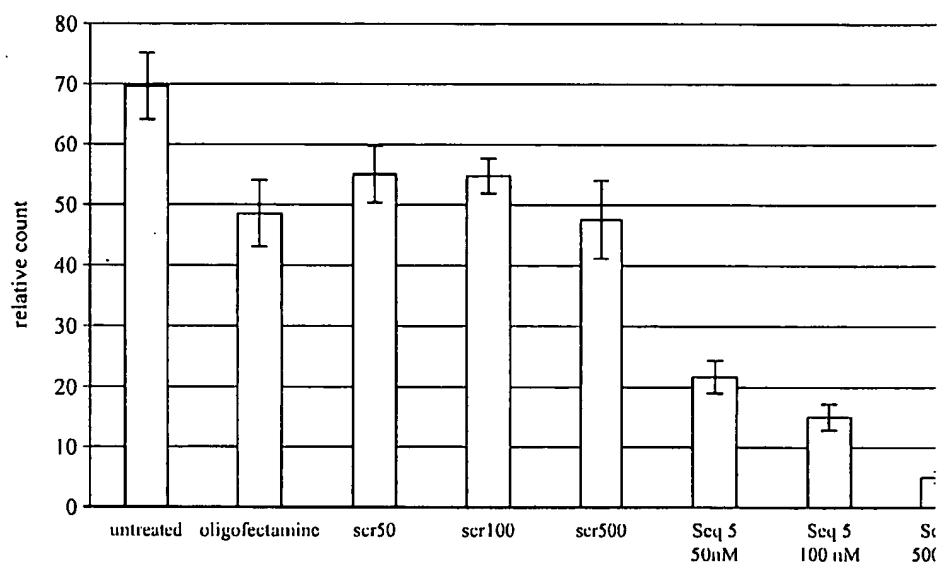


FIG. 3C

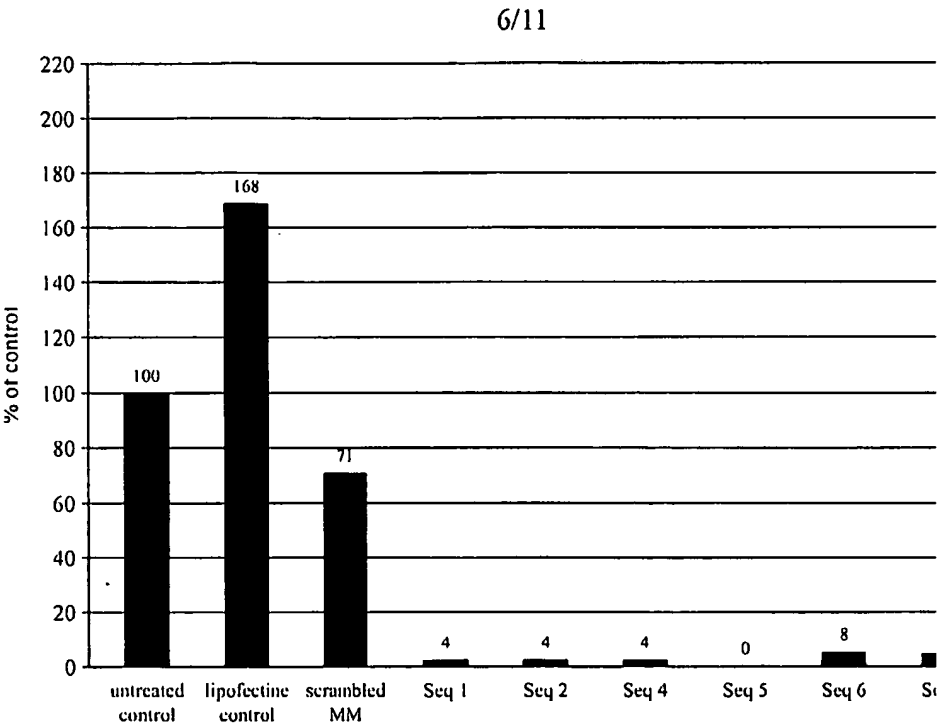


FIG. 3D

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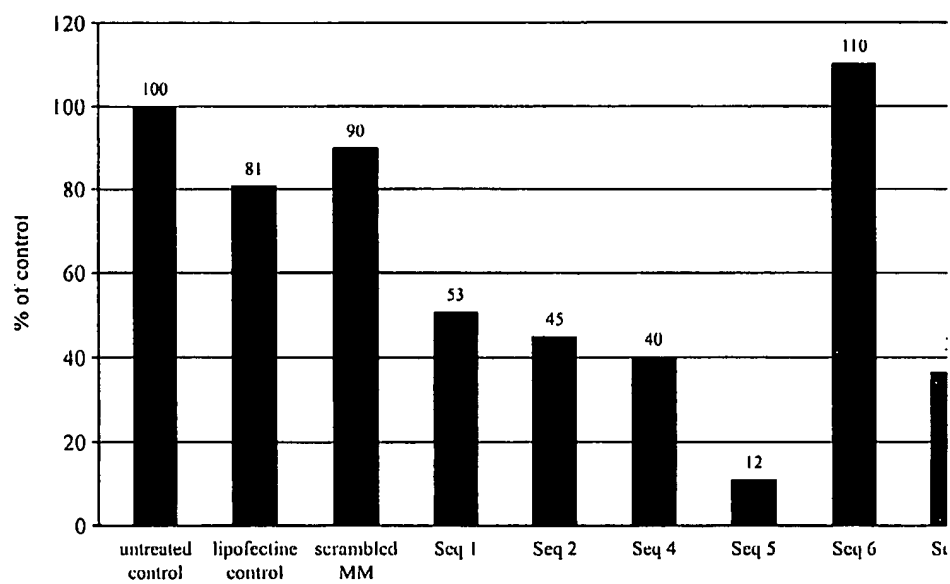


FIG. 3E

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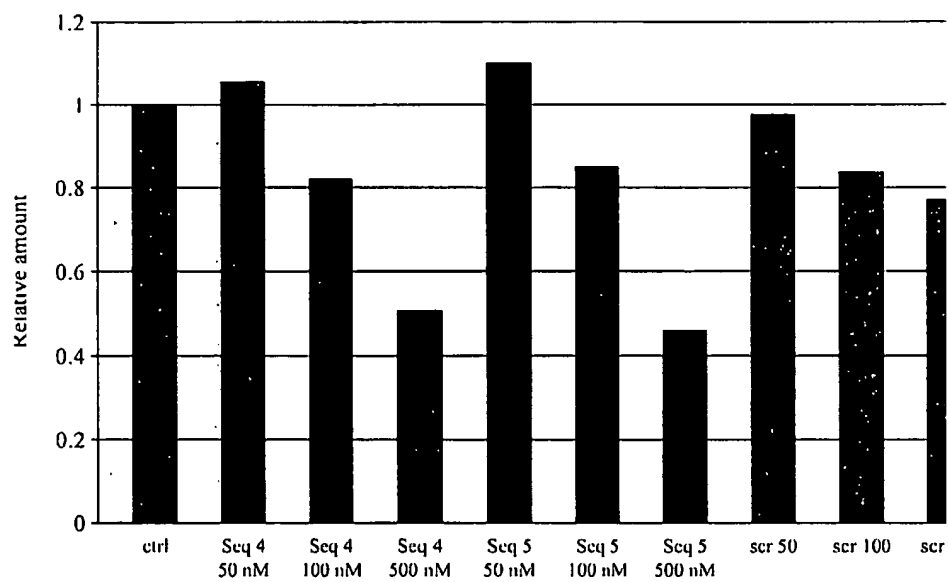


FIG. 4

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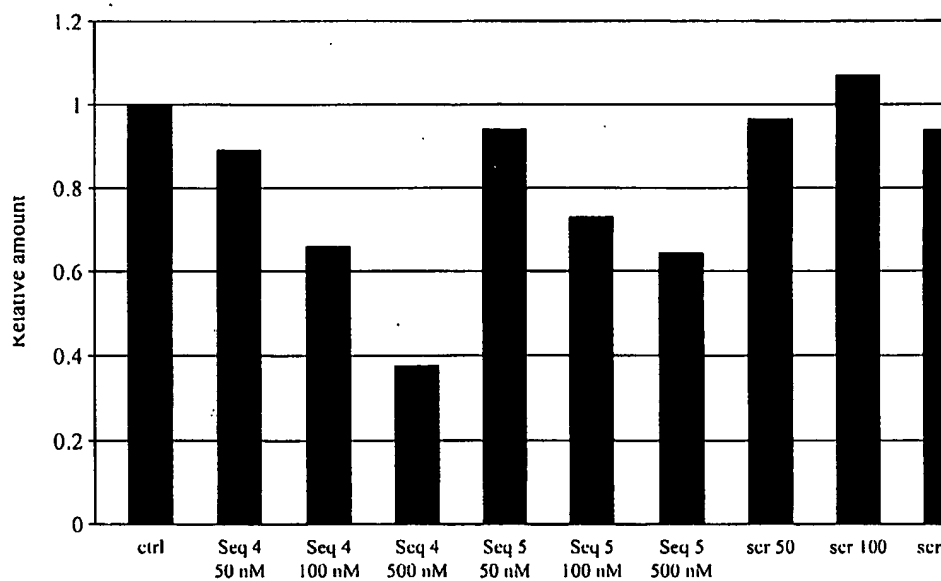


FIG. 5

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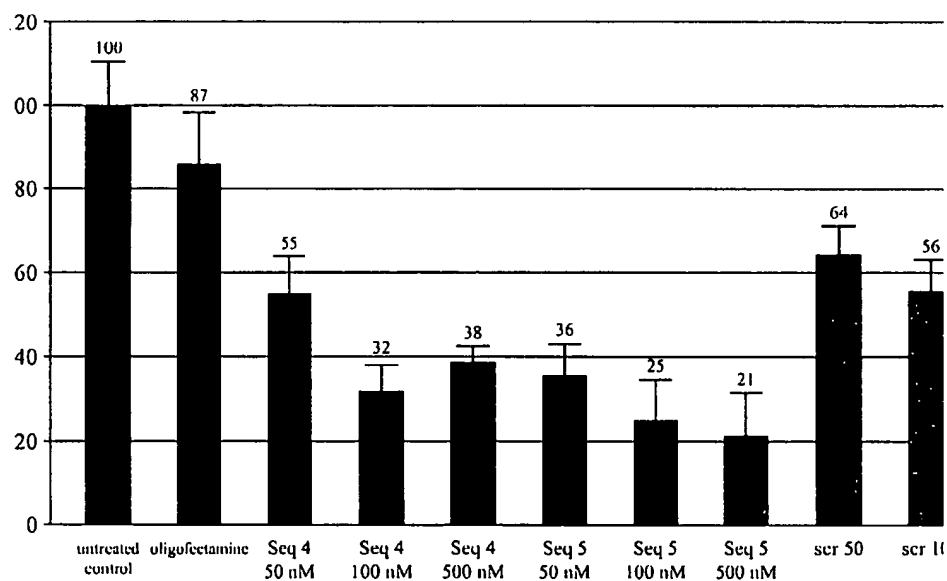


FIG. 6

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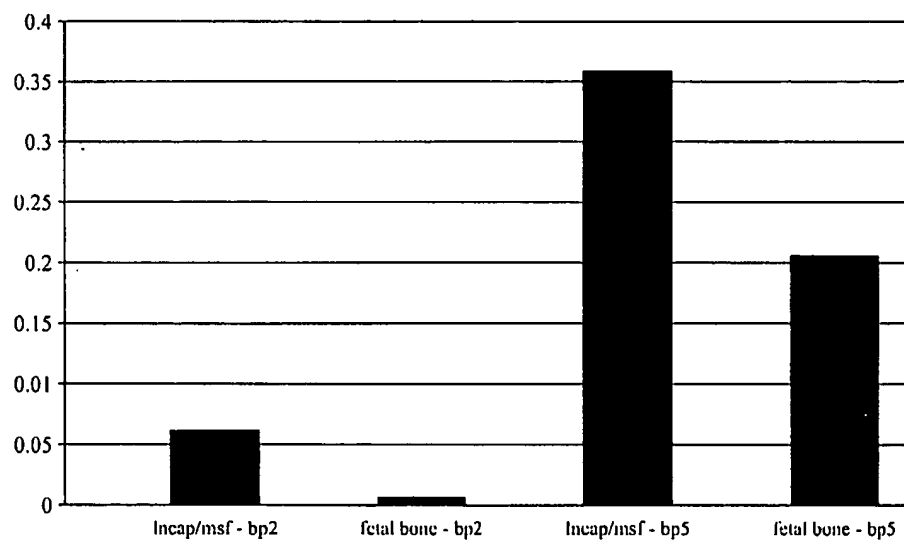


FIG. 7

INTERNATIONAL SEARCH REPORT

International Application No

PCT/CA 03/00061

A. CLASSIFICATION OF SUBJECT MATTER
IPC 7 C12N15/11 A61K31/713

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

MEDLINE, EPO-Internal, EMBASE, BIOSIS, CHEM ABS Data, WPI Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	SHIMASAKI S ET AL: "IDENTIFICATION OF FIVE DIFFERENT INSULIN-LIKE GROWTH FACTOR BINDING PROTEINS (IGFBPS) FROM ADULT RAT SERUM AND MOLECULAR CLONING OF A NOVEL IGFBP-5 IN RAT AND HUMAN" JOURNAL OF BIOLOGICAL CHEMISTRY, AMERICAN SOCIETY OF BIOLOGICAL CHEMISTS, BALTIMORE, MD, US, vol. 266, no. 16, 5 June 1991 (1991-06-05), pages 10646-10653, XP000673586 ISSN: 0021-9258 figure 6 --- -/--	1,5,9



Further documents are listed in the continuation of box C.



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INTERNATIONAL SEARCH REPORT

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PCT/CA 03/00061

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	<p>ZANGEMEISTER-WITKE U ET AL: "A novel bispecific antisense oligonucleotide inhibiting both bcl-2 and bcl-xL expression efficiently induces apoptosis in tumor cells."</p> <p>CLINICAL CANCER RESEARCH: AN OFFICIAL JOURNAL OF THE AMERICAN ASSOCIATION FOR CANCER RESEARCH. UNITED STATES JUN 2000, vol. 6, no. 6, June 2000 (2000-06), pages 2547-2555, XP002241562</p> <p>ISSN: 1078-0432</p> <p>figure 1; table 1</p>	1,5,9
Y	<p>WO 01 05435 A (UNIV BRITISH COLUMBIA ;GLEAVE MARTIN (CA); MIYAKE HIDEAKI (JP))</p> <p>25 January 2001 (2001-01-25)</p> <p>figure 1; examples 1-4</p>	1,5,9
Y	<p>WO 00 69454 A (CHAKRABARTY SHILA ;SINGH POMILA (US); UNIV TEXAS (US))</p> <p>23 November 2000 (2000-11-23)</p> <p>examples 1-5</p>	1,5,9
P,Y	<p>WO 02 22642 A (UNIV BRITISH COLUMBIA ;GLEAVE MARTIN (CA); RENNIE PAUL S (CA); NEL) 21 March 2002 (2002-03-21)</p> <p>examples 1-5</p>	1,5,9
A	<p>FORSYTH E ET AL: "GROWTH INHIBITION OF A HUMAN COLON CANCER CELL LINE BY ANTISENSE OLIGONUCLEOTIDE TO IGFBP-2"</p> <p>GASTROENTEROLOGY, SAUNDERS, PHILADELPHIA, PA,, US,</p> <p>vol. 108, no. 4, 1995, page A726</p> <p>XP002932267</p> <p>ISSN: 0016-5085</p> <p>the whole document</p>	

INTERNATIONAL SEARCH REPORT

International application No.
PCT/CA 03/00061

Box I Observations where certain claims were found unsearchable (Continuation of Item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:

Although claims 6-8 are directed to a method of treatment of the human/animal body, the search has been carried out and based on the alleged effects of the compound/composition.
2. ☐ Claims Nos.:
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of Item 2 of first sheet)

This International Searching Authority found multiple inventions in this International application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/CA 03/00061

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
WO 0105435	A	25-01-2001	AU 6144500 A	05-02-2001
			WO 0105435 A2	25-01-2001
			CA 2375467 A1	25-01-2001
			EP 1200579 A2	02-05-2002
			HU 0201868 A2	28-12-2002
			JP 2003504418 T	04-02-2003
			NO 20020259 A	17-01-2002

WO 0069454	A	23-11-2000	WO 0069454 A1	23-11-2000

WO 0222642	A	21-03-2002	AU 9092001 A	26-03-2002
			WO 0222642 A1	21-03-2002
